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## Amendments to the Claims

This listing of claims replaces all prior versions and listings of claims in the application.

## Listing of Claims

1. (Previously Presented) The use of a conjugate of hydroxyalkylstarch and an allergen in which at least one hydroxyalkylstarch is covalently coupled to the allergen for hyposensitization.

- 2. (Previously Presented) The use as claimed in claim 1, where the hydroxyalkylstarch is coupled directly or via a linker to the allergen.
- 3. (Currently Amended) The use as claimed in claim 1 or 2, where the hydroxyalkylstarch is hydroxyethylstarch, hydroxypropylstarch or hydroxybutylstarch.
- 4. (Currently Amended) The use as claimed in any of claims 1 to 3 claim 1, in which the hydroxyethylstarch has an average molecular weight of from 1 to 300 kDa, preferably an average molecular weight of from 5 to 200 kDa.
- 5. (Currently Amended) The use as claimed in any of the preceding claims claim 1, in which the hydroxyethyl starch has a level of molar substitution of from 0.1 to 0.8 and a C2:C6 substitution ratio in the range from 2 to 20, in each case based on the hydroxyethyl groups.
- 6. (Currently Amended) The use as claimed in any of the preceding claims claim 1, in which the allergen has been selected from the group consisting of polypeptides or proteins.
- 7. (Currently Amended) The use as claimed in any of the preceding claims claim 1, in which the allergen is a glycoprotein.
- 8. (Currently Amended) The use as claimed in any of the preceding claims claim 1, in which the hydroxyalkylstarch is coupled to the polypeptide chain or to one or more of the saccharide chains of the glycoprotein.

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9. (Currently Amended) The use according to any of the preceding claims claim 1 for hyposensitization of allergy sufferers in whom an IgE-mediated sensitization is detected or whose clinical symptoms have been observed.

- 10. (Currently Amended) The use as claimed in any of the preceding claims claim 1, where the specific immunotherapy is employed for the therapy of allergies to pollen, mites, mammalian hair (saliva), fungi, insects, foods and natural rubber/latex.
- 11. (Currently Amended) The use as claimed in any of the preceding claims claim 1, where the therapy is employed for the treatment of asthmatics, hay-fever patients and patients showing other types of clinically relevant reactions to immediate-type allergens.
- 12. (Currently Amended) The use as claimed in any of the preceding claims claim 1, where administration takes place subcutaneously, mucosally, orally, perorally or sublingually.
- 13. (Currently Amended) The use as claimed in any of the preceding claims claim 1, where the immunotherapy is carried out preseasonally or perennially for airborne allergens.
- 14. (Currently Amended) The use any of the preceding claims as claimed in claim 1, where the immunotherapy is carried out for people allergic to insects in the rush or ultra-rush method.
- 15. (New) The use as claimed in any of claims 1 to 3 claim 1, in which the hydroxyethylstarch has an average molecular weight of from 5 to 200 kDa